



Form PTO-1449 (modified)

Atty. Docket No.
UTSD:703USD1Serial No.
10/748,720

List of Patents and Publications for Applicant's

INFORMATION DISCLOSURE STATEMENT

(Use several sheets if necessary)

Applicant

Philip Jordan Thomas, John F. Hunt, William Christian
Wigley and Rhesa D. Stidham

Filing Date:

December 30, 2003

Group:

Unknown

1636

U.S. Patent Documents

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Foreign Patent Documents

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Other Art

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U.S. Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Name	Class	Sub Class	Filing Date of App.
BR	A1	5,120,653	6-9-92	Henderson	435	252.33	10-22-85
BR	A2	6,294,330	9-25-01	Michnick et al.	435	252.3	7-30-98

Foreign Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Country	Class	Sub Class	Translation Yes/No
BR	B1	WO 98/34120	8/6/98	PCT			
BR	B2	WO 98/44350	10/8/98	PCT			

Other Art (Including Author, Title, Date Pertinent Pages, Etc.)

Exam. Init.	Ref. Des.	Citation
BR	C1	Abbas-Terki and Picard, "α-Complemented β-galactosidase. An in vivo model substrate for the molecular chaperone heat-shock protein 90 in yeast," <i>Eur. J. Biochem.</i> , 266:517-523, 1999.
	C2	Betton <i>et al.</i> , "Probing the structural role of an αβ loop of maltose-binding protein by mutagenesis: heat-shock induction by loop variants of the maltose-binding protein that form periplasmic inclusion bodies," <i>J. Mol. Biol.</i> , 262(2):140-150, 1996.
	C3	Blackwell and Horgan, "A novel strategy for production of a highly expressed recombinant protein in an active form," <i>FEBS Lett.</i> , 295:10-12, 1991.
	C4	Blakely <i>et al.</i> , "Epidermal growth factor receptor dimerization monitored in live cells," <i>Nature Biotech.</i> , 18:218-222, 2000.
	C5	Bourot <i>et al.</i> , "Glycine betaine-assisted protein folding in a <i>lysA</i> mutant of <i>Escherichia coli</i> ," <i>J. Biol. Chem.</i> , 275:1050-1056, 2000.
	C6	Brown <i>et al.</i> , "Correcting temperature-sensitive protein folding defects," <i>J. Clin. Invest.</i> , 99:1432-1444, 1997.
	C7	Bruijn <i>et al.</i> , "Aggregation and motor neuron toxicity of an ALS-linked SOD1 mutant independent from wild-type SOD1," <i>Science</i> , 281:1851-1853, 1998.

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Exam. Init.	Ref. Des.	Citation
P5R	C8	Culvenor <i>et al.</i> , "Subcellular localization of the Alzheimer's disease amyloid precursor protein and derived polypeptides expressed in a recombinant yeast system," <i>Amyloid: Int J Exp Clin Invest</i> , 5(2):79-89, 1998.
	C9	Dobson, "Protein misfolding, evolution and disease," <i>TIBS</i> 24:329-332, 1999.
	C10	Johnson and Varshavsky, "Split ubiquitin as a sensor of protein interactions in vivo," <i>Proc Natl Acad Sci U S A</i> , 91(22):10340-4, 1994.
	C11	Foster <i>et al.</i> , "Pharmacological rescue of mutant p53 conformation and function," <i>Science</i> , 286:2507-2510, 1999.
	C12	Harper and Lansbury Jr., "Models of amyloid seeding in Alzheimer's disease and scrapie: mechanistic truths and physiological consequences of the time-dependent solubility of amyloid proteins," <i>Annu. Rev. Biochem.</i> , 66:385-407, 1997.
	C13	Houry <i>et al.</i> , "Identification of <i>in vivo</i> substrates of the chaperonin GroEL," <i>Nature</i> , 402:147-154, 1999.
	C14	Huang <i>et al.</i> , "NMR structure and mutagenesis of the Fas (APO-1/CD95) death domain," <i>Nature</i> , 384:638-641, 1996.
	C15	Hung <i>et al.</i> , "Crystal structure of the ATP-binding subunit of an ABC transporter," <i>Nature</i> , 396:703-707, 1998.
	C16	Huth <i>et al.</i> , "Design of an expression system for detecting folded protein domains and mapping macromolecular interactions by NMR," <i>Protein Sci.</i> , 6:2359-2364, 1997.
	C17	Johnsson and Varshavsky, "Split ubiquitin as a sensor of protein interactions in vivo," <i>Proc. Natl. Acad. Sci. USA</i> , 91:10340-10344, 1994. OF RECORD - C10
	C18	Kapust and Waugh, " <i>Escherichia coli</i> maltose-binding protein is uncommonly effective at promoting the solubility of polypeptides to which it is fused," <i>Protein Science</i> , 8:1668-1674, 1999.
	C19	King and Sorscher, "Recombinant synthesis of cystic fibrosis transmembrane conductance regulator and functional nucleotide-binding domains," <i>Methods Enzymol.</i> , 292:686-697, 1998.
✓	C20	Ko <i>et al.</i> , "The cystic fibrosis transmembrane conductance regulator," <i>J. Biol. Chem.</i> , 268:24330-24338, 1993.

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P50	C21	Lee <i>et al.</i> , "Effect of the N-terminal hydrophobic sequence of hepatitis B virus surface antigen on the folding and assembly of hybrid β -galactosidase in <i>Escherichia coli</i> ," <i>Eur. J. Biochem.</i> , 187:417-424, 1990.
	C22	Luzzago and Cesareni, "Isolation of point mutations that affect the folding of the H chain of human ferritin in <i>E. coli</i> ," <i>EMBO J</i> , 8:569-576, 1989.
	C23	Maxwell <i>et al.</i> , "A simple <i>in vivo</i> assay for increased protein solubility," <i>Protein Science</i> , 8:1908-1911, 1999.
	C24	Nixon and Benkovic, "Improvement in the efficiency of formyl transfer of a GAR transformylase hybrid enzyme," <i>Protein Engineering</i> , 13(5):323-327, 2000.
	C25	Opal and Paulson, "Genetic instabilities and hereditary neurological diseases," <i>Am J. Hum. Genet.</i> , 63(6):1921, 1998.
	C26	Papouchado <i>et al.</i> , "Expression of properly folded human glutamate decarboxylase 65 as a fusion protein in <i>Escherichia coli</i> ," <i>Eur. J. Biochem.</i> , 246:350-359, 1997.
	C27	Pelletier <i>et al.</i> , "An <i>in vivo</i> library-versus-library selection of optimized protein-protein interactions," <i>Nature Biotech.</i> , 17:683-690, 1999.
	C28	Qu and Thomas, "Alteration of the cystic fibrosis transmembrane conductance regulator folding pathway," <i>J. Biol. Chem.</i> , 271(13):7261-7264, 1996.
	C29	Rao <i>et al.</i> , "Rhodopsin mutation G90D and a molecular mechanism for congenital night blindness," <i>Nature</i> , 367:639-642, 1994.
	C30	Sugihara and Baldwin, "Effects of 3' end deletions from <i>Vibrio hrvveyi</i> luxB gene on luciferase subunit folding and enzyme assembly: generation of temperature-sensitive polypeptide folding mutants," <i>Biochemistry</i> , 27:2872-2880, 1988.
	C31	Tan and Pepys, "Amyloidosis," <i>Histopathology</i> , 25:403-414, 1994.
	C32	Thomas <i>et al.</i> , "Altered protein folding may be the molecular basis of most cases of cystic fibrosis," <i>FEBS Lett.</i> , 312:7-9, 1992.
	C33	Thomas <i>et al.</i> , "Defective protein folding as a basis of human disease," <i>TIBS</i> , 20:456-459, 1995.
✓	C34	Valois <i>et al.</i> , "Utilisation of the PCA strategy to study the folding of the RBD of raf," <i>FASEB Journal</i> , 13:A1387, 330, 1999.

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Exam. Init.	Ref. Des.	Citation
FR	C35	Waldo <i>et al.</i> , "Rapid protein-folding assay using green fluorescent protein," <i>Nature Biotechnology</i> , 17:691-695, 1999.
J	C36	Wang <i>et al.</i> , "Expression and purification of the first nucleotide-binding domain and linker region of human multidrug resistance gene product: comparison of fusions to glutathione S-transferase, thioredoxin and maltose-binding protein," <i>Biochem J.</i> , 338:77-81, 1999.
W	C37	Wood <i>et al.</i> , "Prolines and amyloidogenicity in fragments of the Alzheimer's peptide β /A4," <i>Biochemistry</i> , 34(3):724-730, 1995.

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